



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

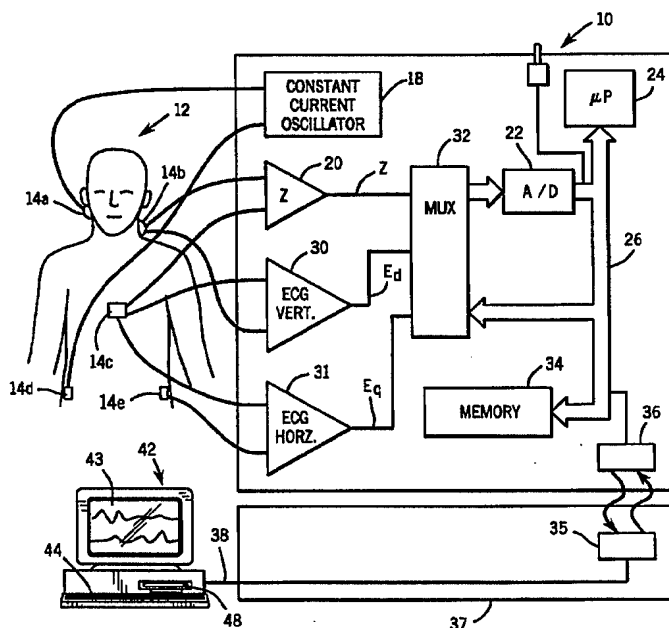
(51) International Patent Classification ⁶ : A61B 5/05, 5/029	A1	(11) International Publication Number: WO 96/01586 (43) International Publication Date: 25 January 1996 (25.01.96)
--	-----------	--

(21) International Application Number: PCT/US95/08856

(22) International Filing Date: 23 June 1995 (23.06.95)

(30) Priority Data:
08/271,689 7 July 1994 (07.07.94) US(71) Applicant: REINING INTERNATIONAL LTD. [US/US];
5930 Seminole Center Court, Madison, WI 53611 (US).(72) Inventor: REINING, William, N.; 4580 Garfoot Road, Cross
Plains, WI 53523 (US).(74) Agents: BAXTER, Keith, M. et al.; Quarles & Brady, Suite
2550, 411 East Wisconsin Avenue, Milwaukee, WI 53202-
4497 (US).(81) Designated States: European patent (AT, BE, CH, DE, DK,
ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).**Published***With international search report.**Before the expiration of the time limit for amending the
claims and to be republished in the event of the receipt of
amendments.*

(54) Title: IMPEDANCE CARDIOGRAPH APPARATUS AND METHOD



(57) Abstract

An impedance cardiograph (10) which determines cardiac output from a measurement of variations of chest impedance (2), provides method of calculating the effect of patient volume on the measurement from measurable patient height (H) and chest circumference (C), and provides a correction process that identifies as a source of error variations in the first derivative of impedance. This latter source of error is minimized by preprocessing the impedance derivative value with a compression function which reduces the range of values of the impedance derivative when that value differs significantly from a norm of the population.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgyzstan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

IMPEDANCE CARDIOGRAPH APPARATUS AND METHOD

Field of the Invention

The present invention relates to impedance cardiographs which determine cardiac output by evaluating changes in impedance across the patient's chest cavity.

5 Background of the Invention

Impedance cardiography is a non-invasive technique for determining cardiac performance in humans. When such equipment is employed, a high frequency electric signal is applied to the patient across outer electrodes
10 positioned, for example, on the patient's head and lower thorax. Voltage differences between sensing inner electrodes positioned between the outer electrodes on the patient's neck and chest are measured and used to compute an impedance (Z). The impedance is based on the low
15 magnitude, known electrical current passing between the outer electrodes.

In 1932, Atzler and Leyman reported that cardiac output of a human could be determined by such impedance methods in *Über ein neues Verfahren zur Darstellung der*
20 *Herztätigkeit (Dielektrographie)*, *Arbeitsphysiologie*, 5:636-680. In 1966, Kubicek reported the ability to correlate changes in base line impedance and the first derivative of impedance to stroke volume (SV) according to the following Equation (1) disclosed in U.S. Patent
25 3,340,867 and in the publication: *Development and*

Evaluation of an Impedance Cardiac Output System,
Aerospace Medicine, 37:1208-1212.

$$SV = \frac{\rho * L^2 * \frac{d\hat{Z}}{dt} * L_{vet}}{Z_0^2} \quad (1)$$

where:

5 ρ is the resistivity of blood;

L is the spacing between the sensing electrodes;

Z_0 is an average or baseline impedance; and

$\frac{d\hat{Z}}{dt}$ is the magnitude of the peak negative value of

the time derivative of the impedance Z for a period of
10 time, typically a second.

Cardiac output may be deduced from stroke volume by multiplying the latter times the heart rate.

Although the Kubicek formula provides a value that correlated with cardiac output, the absolute accuracy of
15 the method remained doubtful and, in particular, subjects with certain cardiovascular problems show values with great inaccuracies.

In 1982, Sramek proposed a modification of the Kubicek formula of equation (1) which resolved the base
20 line impedance Z_0^2 into a dynamic and static component as reported in the publication: *Cardiac Output by Electrical Impedance*, Med. Elect., 2:274-290. The static term Z_{0s} was described by the following equation:

$$Z_{0s} = \frac{\rho * L}{A} \quad (2)$$

25 where A is the area of the thorax being measured.

The dynamic component Z_{0d} was simply the baseline or average of the impedance being measured:

$$Z_0 d = Z_0 \quad (3)$$

Incorporating the static term and dynamic term into the Kubicek equation provides the following formula:

$$SV = \frac{A * L * \frac{d\hat{Z}}{dt} * L_{vet}}{Z_0} \quad (4)$$

5 The value of A may be estimated by approximating the chest as a cylinder in which case equation (4) becomes:

$$SV = \frac{C^2 * L * \frac{d\hat{Z}}{dt} * L_{vet}}{4 * \pi * Z_0} \quad (5)$$

where C is circumference of the chest near the area of measurement. Alternatively, Sramek proposed that the
10 term

$\frac{C^2 * L}{4 * \pi}$ be replaced with either $\frac{L^3}{4.25}$, or $\frac{(1.7 * H)^3}{4.25}$ where H is the height of the patient because $1.7 * H$ approximates L.

These approximations did not produce good results and so Sramek was ultimately led to produce a set of
15 charts attempting to establish correlation between area and the three factors of gender, height and weight.

In 1986, Bernstein proposed a modified equation in which the separation of the electrodes and the height of the patient were considered, in the following form:

$$20 \quad SV = \frac{\left(\frac{L + 1.5 * H}{2} \right)^3 * \frac{d\hat{Z}}{dt} * L_{vet}}{4.25 * Z_0} \quad (6)$$

All of the above methods suffer from lack of accuracy and indicate, in some subjects, falsely high or low values of stroke volume.

Summary of the Invention

The present invention provides an improved method and apparatus for deducing stroke volume (and hence cardiac output) from impedance measurements. The

5 invention provides an improved estimation of body volume and a processing of the derivative of the impedance signal that improves the reliability of the derived values of stroke volume and cardiac output.

Specifically, in an impedance cardiograph for use on
10 a human patient having a height of H and a chest circumference C, a means for applying an electrical excitation signal to the patient is used in conjunction with electrodes positioned on the chest with a separation distance of L, the electrodes producing a first
15 electrical impedance signal Z which varies with impedance changes in the patient. A user input device, such as a keyboard, is provided to enter data on the height, electrode separation distance and circumference to produce corresponding second and third electrical signals
20 H, L and C indicating those values. An electrical circuit, which may be an electronic computer, receives the second and third electrical signal and provides an indication of the patient's cardiac stroke volume, SV, as a function of Z, A, L and left ventricular ejection
25 interval L_{vet} where A is deduced by the following approximation:

$$A = \frac{L}{K \cdot H \cdot C}$$

where K is a predetermined constant.

Thus, the impedance cardiograph may calculate SV according to the following formula:

$$SV = \frac{L}{K \cdot H \cdot C} \frac{\rho \cdot C^2 \cdot L \cdot L_{vet}}{4 \cdot \pi \cdot Z_0} \frac{d\hat{Z}'}{dt}$$

which may be simplified to:

5
$$SV = \frac{\rho \cdot C^2 \cdot L^2 \cdot L_{vet}}{4 \cdot K \cdot H \cdot \pi \cdot Z_0} \frac{d\hat{Z}'}{dt}$$

or by combining constants:

$$SV = \frac{\rho \cdot C^2 \cdot L^2 \cdot L_{vet}}{K \cdot H \cdot Z_0} \frac{d\hat{Z}'}{dt}$$

Thus, it is one object of the invention to provide a simple, yet more accurate characterization of a critical
 10 term A used in the calculation of stroke volume from patient impedance. In the present invention, the area of the impedance measurement, which is difficult to measure, is accurately derived from the readily measured values of chest circumference and patient height.

15 The present invention has also recognized that variation in the derivative of impedance, one of the factors used in deducing stroke volume, is a significant source of inaccuracy in the computed stroke volume. Accordingly, whereas $\frac{d\hat{Z}'}{dt}$ may simply be the magnitude of
 20 the minimum derivative of impedance with time, it may also be compressed to reduce the amount that this maximum deviates from the norm. One method of weighting the maximum is according to the formula:

$$\frac{d\hat{Z}'}{dt} = \frac{d\hat{Z}_n}{dt} = \frac{\frac{d\hat{Z}}{dt}}{\sqrt{\frac{d\hat{Z}}{dt} / \frac{d\bar{Z}}{dt}}}$$

where:

$\frac{d\hat{Z}}{dt}$ is the magnitude of the minimum time derivative of Z;

5 $\frac{d\hat{Z}_n}{dt}$ is the compressed value of $\frac{d\hat{Z}}{dt}$;

$\frac{d\bar{Z}}{dt}$ is a predetermined normal value and is an average value of $\frac{d\hat{Z}}{dt}$ for a population.

Thus, it is yet another object of the invention to reduce the effect of a significant source of error in the calculation of stroke volume from impedance by
10 implementation of a normal based weighting system.

The foregoing and other objects and advantages of the invention will appear from the following description. In the description, reference is made to the accompanying
15 drawings which form a part hereof and in which there is shown by way of illustration, a preferred embodiment of the invention. Such embodiment does not necessarily represent the full scope of the invention, however, and reference must be made therefore to the claims herein for
20 interpreting the scope of the invention.

Brief Description of the Drawings

Fig. 1 is a block diagram of the circuitry for an impedance cardiograph according to the present invention, showing a computer as used to analyze the impedance data
25 to produce a value of stroke volume and cardiac output;

and

Fig. 2 is a flow chart of the software employed by the computer of Fig. 1 in analyzing the impedance and ECG signals acquired from the patient.

5 Detailed Description of the Preferred Embodiment

Referring to Fig. 1, a portable impedance cardiograph 10 is connected to a patient 12 by five patch electrodes 14(a)-(e). The first electrode 14(a) may be positioned on the patient's skin behind the right ear at the level of the ear canal. The second electrode 14(b) may be located at the left side of the neck on a flat surface approximately between the level of the chin and base of the hairline at least 5 cm below the level of electrode 14(a). The third electrode 14(c) may be located just above the base of the sternum on the anterior median lines. The fourth and fifth electrodes 14(d) and 14(e) may be located at least 5 cm below the electrode 14(c) on the patient's right and left sides respectively at the costal arch and the anterior axillary line.

Electrodes 14(a) and (d) are attached to an oscillator 18 which produces a constant current of approximately 1 milliamperes RMS through the patient 12. This electrical excitation establishes a series of equal potential surfaces through the patient 12 perpendicular to a line extending between the two outer electrodes 14(a) and 14(d).

Electrodes 14(b) and 14(c) may sense the equal

potential lines generated by the current flowing between the outer electrodes 14(a) and 14(d). Because the current between the outer electrodes 14(a) and 14(d) is of constant amplitude, the amplitude of the voltage
5 sensed between the inner electrodes 14(b) and 14(c) is proportional to the thoracic impedance of the patient 12. The inner electrodes 14(b) and 14(c) are connected to differential amplifier 20 producing a signal Z.

The amplifier 20 includes isolation circuitry that
10 electrically isolates the inner electrodes 14(b) and 14(c) from the subsequent circuitry. Differential amplifier 20 also includes a precision half wave rectifier and low pass filter so as to provide a slowly varying DC signal whose value is proportional to the
15 impedance being measured. The input impedance of the differential amplifier 20 is very high (e.g. 10 megohms) as compared to the impedance of the patient 12 between the inner electrodes 14(b) and 14(c). Thus, negligible current will flow through the inner electrodes 14(b) and
20 14(c) to amplifier 20.

The impedance signal Z is received by a multiplexer 32, such as are known in the art, to be periodically connected to an analog to digital converter 22 which samples the signal and provides a binary data word that
25 may be read by microprocessor 24 via a bus 26.

A first, vertical ECG signal is measured across electrodes 14(b) and 14(c) lying generally along a generally vertical line. This ECG signal is received by differential amplifier 30 to produce an

electrocardiograph signal E_v according to techniques well known in the art.

A second, horizontal ECG signal is measured across electrodes 14(c) and 14(e) lying generally along a horizontal line. This ECG signal is received by differential amplifier 31 to produce an electrocardiograph signal E_h . Both signals E_v and E_h are received by multiplexer 32 which periodically connects these signals to the analog to digital converter 22 to be sampled and converted to digital words for transmission on internal bus 26.

Differential amplifiers 30 and 31 receiving the ECG signal also include isolation circuitry and a low pass filter having a cut-off frequency such as to substantially remove the 50 kHz oscillator signal from oscillator 18. As will be described further below, the direct and quadrature ECG signals are combined to produce a single ECG signal largely independent of the electrical orientation of the patient's heart.

Also attached to bus 26 is computer memory 34 which may be composed of both random access memory ("RAM") and read only memory ("ROM") according to well known computer architectures. Memory 34 provides a means for storage of the binary representations of signals Z and E_v and E_h under the control of microprocessor 24, and also holds a stored program defining the operation of the microprocessor 24 for the calculation of cardiac output as will be described.

Also attached to bus 26 is a pushbutton switch 25

which may be used by the patient to mark the occurrence of some event, such as a cardiac episode, during the recording of data from the patient 12 as will be described.

5 The bus 26 also communicates with an infra-red transceiver 36, which permits the microprocessor to transmit and receive data to and from a similar transceiver 35 in a detachable base unit 37. The transceiver 35 is connected by a serial cable 38 to a
10 desk-top computer 42 having a video monitor 43, a keyboard 44 and a disk drive 48 such as are known in the art. As such the impedance cardiograph 10 is portable and may be powered by internal batteries so as to be carried with the patient in the manner of a Holter
15 monitor.

 Generally, the impedance cardiograph receives signals from the patient 12, isolates, amplifies and filters those signals, and then translates the signals to digital values which may be read and stored by
20 microprocessor 24 to be processed according to the stored program in memory 34. Results of the processing may be transmitted to the computer 42 to be displayed on the video monitor 43 or saved on disk 48.

 The operation of the impedance cardiograph 10
25 according to the stored program is controlled by a human operator through keyboard 44. The operator prepares the patient 12 for the impedance measurements and may enter certain data to the keyboard 44 that characterizes the patient 12 and that is necessary for the analysis of the

signals from the patients 12 as will be described. This analyses is done in partially in the microprocessor 24 so as to reduce the amount of data to be stored in memory 34, but, as will be understood in the art, the analyses of the data may be shared between the microprocessor 24 and the computer 42 as a matter of engineering choice.

Referring now to Fig. 2, at the first step in the analyses, indicated by process block 50, certain data related to the particular patient 12 or related to fundamental and essentially universal physiological parameters, may be entered by the operator.

The first of these parameters is ρ which is the resistivity of blood in ohms-cm. Generally, this value may be approximated as a constant for all patients, however, it may be modified by the operator in abnormal cases based on the measurement of hematacrit.

A second value, C , is the circumference of the patient's chest in cm taken at the site of inner electrode 14(c) around the patient's chest.

The value L is also input, being the distance between the inner electrodes 14(b) and 14(c) in cm. The height of the patient, H , in cm is also entered.

Once the necessary fixed parameters are entered at process block 50, two variable parameters: heart rate HR and ventricular ejection time L_{vet} are entered per process block 52. For this purpose, the ECG signal E may be directly displayed on the video monitor 43 so that these quantities may be determined according to methods well known in the art. The ECG signal is calculated from

the vector sum of the values of E_v and E_h most simply as follows:

$$ECG = \sqrt{E_v^2 + E_h^2}$$

This vector summing reduces the need to precisely
5 orient the ECG electrodes with respect to an electrical
polarity of the heart and therefore in practice provides
a superior ECG signal.

Generally, L_{vet} is the time between the opening of
the aortic valve and the closing of the aortic valve.
10 The heart rate is simply the number of beats per second
which is the inverse of the period between successive R
waves. The heart rate may be averaged over a number of
beats according to methods well known in the art. Both
quantities may be determined by inspection by the
15 operator or preferably may be determined automatically
after sufficient ECG and Z data is acquired as indicated
by process block 52. The value of L_{vet} in the preferred
embodiment is determined by analyzing the impedance
signal Z to measure a period beginning when $\frac{dZ}{dt}$ is first
20 less than zero and ending when the value of $\frac{dZ}{dt}$ reaches a
local maximum above zero. The heart rate HR is measured
by detecting and counting R waves in the ECG signal.

The acquisition of the ECG and Z signal per process
block 54 continues. This acquisition is on a continuous
25 basis and occurs concurrently with the subsequent
calculations so that the cardiac output may be
continuously stored in memory 34 or displayed in
essentially a real-time manner.

The acquired impedance data is in the form of discrete samples taken approximately 300 times per second, each sample which may be represented by Z_i where i is an index number of the particular sample. As each sample Z_i is acquired, it is stored in consecutive addresses in memory 34 to indicate its relative position with respect to other samples and to indicate the time of the sample indirectly through the constant sampling rate.

Because the impedance cardiograph 10 is portable, there is a risk that artifacts may be introduced into the impedance measurement by electrode movement. This is because the measured impedance values are approximately two orders of magnitude lower than the electrode to skin resistance. Accordingly, the impedance data over a period of approximately one minute is "ensemble" averaged, per process block 56, to reduce its noise content. Ensemble averaging is a well known technique in which blocks of impedance data are averaged on a point-by-point basis with other blocks of impedance data so that the averaged points are from corresponding portions of the impedance waveform cycle. Thus, the shape of the impedance waveform is not destroyed in the averaging process. In order to perform such ensemble averaging, it is necessary to identify a common fiducial point to align the blocks of data. Selection of the fiducial point must be extremely precise, otherwise the characteristics of the impedance waveform will be "blurred" by a mis-registration of other blocks.

This fiducial point may be the peak of the R wave of

the ECG signal. Normal techniques for determining the time of the R wave, such as may be used for the measurement of heart rate, however, are not suitably accurate for the purpose of ensemble averaging.

5 Accordingly an extremely accurate identification process is used. First, as represented by process block 58, the ECG signal is monitored to isolate a standard R wave. Only portions of the received ECG signal having no detectable artifacts or noise are considered. This
10 standard R wave is then correlated to the incoming ECG signal to identify the precise location of the R wave (by the value of highest correlation). This location is used at the point of common alignment for the impedance waveforms to be ensemble averaged. Periodically, a new
15 standard R wave is obtained so that the standard remains current over time.

After pairs of data Z_i and Z_{i+1} are acquired and averaged, a derivative value is $\frac{dZ_i}{dt}$ may be computed by a simple subtraction of adjacent samples of the ensemble
20 average per process block 58, that is:

$$\frac{dZ_i}{dt} = Z_{i+1} - Z_i \quad (7)$$

Alternatively, in order to reduce the presence of 50 Hz or 60 Hz noise, this derivative computation can employ samples $Z_{i+6} - Z_i$ or Z_{i+5} , respectively.

25 It has been determined that variation in the magnitude of the minimum of this value, $\frac{d\hat{Z}}{dt}$, is a significant source of error in the calculation of cardiac output. Accordingly, at process block 60, a compressed

derivative, $\frac{d\hat{Z}_n}{dt}$ is computed according to the following formula:

$$\frac{d\hat{Z}_n}{dt} = \frac{\frac{d\hat{Z}}{dt}}{\sqrt{\frac{d\hat{Z}}{dt} / \frac{d\bar{Z}}{dt}}} \quad (8)$$

where:

5 $\frac{d\hat{Z}}{dt}$ is the magnitude of the minimum time derivative

of Z as previously defined;

$\frac{d\hat{Z}_n}{dt}$ is the compressed value of $\frac{d\hat{Z}}{dt}$;

$\frac{d\bar{Z}}{dt}$ is the predetermined normal maximum value and is an average value $\frac{d\hat{Z}}{dt}$ for a population and is about 1.73

10 ohms per second.

Equation (8) has the effect of reducing the excursions of $\frac{d\hat{Z}}{dt}$ from $\frac{d\bar{Z}}{dt}$. Other nonlinear compression systems may also be used provided they have the effect of compressing $\frac{d\hat{Z}}{dt}$ about the norm.

15 At succeeding process block 52, the value of Z_0 is also computed. Z_0 is the base transthoratic impedance and in this implementation, simply the average value of Z for one cardiac cycle. Because of the need to average a number of samples, when the sampling is first begun, no

20 display is provided to the video terminal 43 until sufficient samples have been made to insure the accuracy of this value Z_0 .

At process block 64 a stroke volume may be calculated according to the following formula:

$$SV = \frac{L}{K \cdot H \cdot C} \frac{\rho \cdot C^2 \cdot L \cdot L_{vet}}{4 \cdot \pi \cdot Z_0} \frac{d\hat{z}}{dt} \quad (9)$$

It will be recognized that this is simply Equation (5) with the addition of a factor ρ and the addition of a factor $\frac{L}{K \cdot H \cdot C}$, this latter factor approximating a slice of

5 body volume in the area of the impedance measurement.

This calculation of stroke volume has shown significant improvements in accuracy and correlation coefficients in clinical studies.

Also at process block 64, cardiac output may be
10 determined by multiplying the stroke volume times the heart rate:

$$CO = SV \cdot HR \quad (10)$$

As cardiac output is computed, it is displayed in graphical form on video monitor 43. Thus, the operator
15 is provided with concurrent ECG data and cardiac output data on an essentially real-time basis per the display indicated by process block 66.

After each updating of the display of 43 is accomplished, the program acquire additional data until
20 the measurement session is complete.

While this invention has been described with reference to particular embodiments and examples, other modifications and variations will occur to those skilled in the art in view of the above teachings. Accordingly,
25 the present invention is not limited to the preferred embodiment described herein, but is instead defined in the following claims.

Claims

I claim:

1. An impedance cardiograph used to evaluate cardiac output of a human patient having a height and a chest circumference and chest cross-sectional area A comprising:

5 means for applying an electrical excitation signal to the chest of a patient;

electrodes, adapted to be positioned on the patient with a separation distance of L thereby being responsive to the excitation signal, for producing a first
10 electrical signal Z which varies with impedance changes in the patient;

input device for receiving values of the height and circumference and providing corresponding second and third electrical signals H and C indicating those values;

15 electrical circuit means for receiving the first, second and third electrical signal and providing an indication of the patient's cardiac stroke volume SV as a function of Z, A, and L;

where A is deduced by the following approximation:

20
$$\frac{L}{K \cdot H \cdot C}$$

where K is a predetermined constant.

2. The impedance cardiograph of claim 1 wherein the electric circuit means calculates SV according to the following formula:

$$SV = \frac{L}{K \cdot H \cdot C} \frac{\rho \cdot C^2 \cdot L \cdot L_{vet}}{4 \cdot \pi \cdot Z_0} \frac{d\hat{Z}'}{dt}$$

5 where

ρ is the resistivity of blood;

Z_0 is an average baseline impedance;

L_{vet} is a left ventricular ejection interval; and

$\frac{d\hat{Z}'}{dt}$ is a function of the time derivative of the

10 impedance Z.

3. The impedance cardiograph of claim 2 wherein $\frac{d\hat{Z}'}{dt}$ is the magnitude of the minimum time derivative of Z.

4. The impedance cardiograph of claim 2 wherein $\frac{d\hat{Z}'}{dt}$ is a compressed value of a time derivative of Z about a predetermined normal value so that $\frac{d\hat{Z}'}{dt}$ has a value less than the time derivative of Z if the time derivative of Z is greater than a predetermined normal value and $\frac{d\hat{Z}'}{dt}$ has a value greater than the time derivative of Z if the time derivative of Z is less than the predetermined normal value.

5. The impedance cardiograph of claim 1 wherein K is substantially 0.875.

6. An impedance cardiograph used to evaluate cardiac output of a human patient comprising:

means for applying an electrical excitation signal to the chest of a patient;

5 electrodes, adapted to be positioned on the patient for producing a first electrical signal Z which varies with impedance changes in the patient;

a non-linear processor for calculating a time derivative of Z, and providing a compressed derivative,
10 having a value less than the time derivative of Z if the time derivative of Z is greater than a predetermined normal value and having a value greater than the time derivative of Z if the time derivative of Z is less than the predetermined normal value;

15 electrical circuit means for providing an indication of patient's cardiac stroke volume SV as a function of the compressed time derivative.

7. The impedance cardiograph of claim 6 wherein the non-linear processor produces the compressed value of the time derivative of Z according to the following formula:

$$\frac{d\hat{Z}_n}{dt} = \frac{\frac{d\hat{Z}}{dt}}{\sqrt{\frac{d\hat{Z}}{dt} / \frac{d\bar{Z}}{dt}}}$$

5

where:

$\frac{d\hat{Z}}{dt}$ is a time derivative of Z

$\frac{d\hat{Z}_n}{dt}$ is the compressed time derivative of Z

$\frac{d\bar{Z}}{dt}$ is the predetermined normal value and is an average value of $\frac{d\hat{Z}}{dt}$ for a population.

8. The impedance cardiograph of claim 6 wherein the predetermined normal value is substantially 1.73 ohms per second.

9. A method of evaluating the cardiac output of a human patient having a height H and a chest circumference C and chest cross-sectional area A comprising the step of:

5 (a) applying an electrical excitation signal to the chest of a patient;

(b) positioning electrodes on the patient with a separation distance of L thereby being responsive to the excitation signal to produce a first electrical signal Z
10 which varies with impedance changes in the patient;

(c) employing an electronic computer to:

(i) receive values of the height and circumference and providing corresponding second and third electrical signals H and C indicating those values;
15 (ii) evaluate the patient's cardiac stroke volume SV as a function of Z, A, L and left ventricular ejection interval L_{vet} ;

where A is deduced by the following approximation:

$$\frac{L}{K \cdot H \cdot C}$$

20 in which K is a predetermined constant.

10. The method claim 9 wherein the electronic computer calculates SV according to the following formula:

$$SV = \frac{L}{K \cdot H \cdot C} \frac{\rho \cdot C^2 \cdot L \cdot L_{vet}}{4 \cdot \pi \cdot Z_0} \frac{d\hat{Z}'}{dt}$$

5

where

ρ is the resistivity of blood;

Z_0 is an average baseline impedance; and

$\frac{d\hat{Z}'}{dt}$ is a function of the time derivative of the

impedance Z.

11. The method of claim 10 wherein $\frac{d\hat{Z}'}{dt}$ is the magnitude of the minimum time derivative of Z.

12. The method of claim 10 wherein $\frac{d\hat{Z}'}{dt}$ is a compressed value of the time derivative of Z about a predetermined normal value so that $\frac{d\hat{Z}'}{dt}$ has a value less than the time derivative of Z if the time derivative of Z is greater than a predetermined normal value and $\frac{d\hat{Z}'}{dt}$ has a value greater than the time derivative of Z if the time derivative of Z is less than the predetermined normal value.

13. The method of claim 12 wherein K is substantially 0.875.

14. A method of evaluating cardiac output of a human patient comprising the steps of:

(a) applying an electrical excitation signal to the chest of a patient;

5 (b) positioning electrodes on the patient for producing a first electrical signal Z which varies with impedance changes in the patient;

(c) determining a compressed derivative of the impedance signal Z, having a value less than the time derivative of Z if the time derivative of Z is greater
10 than a predetermined normal value and having a value greater than the time derivative of Z if the time derivative of Z is less than the predetermined normal value; and

15 (e) evaluating the patient's cardiac stroke volume SV as a function of the compressed time derivative.

15. The method of claim 14 step (c) is according to the following formula:

$$\frac{\hat{dZ}_n}{dt} = \frac{\frac{\hat{dZ}}{dt}}{\sqrt{\frac{\hat{dZ}}{dt} / \frac{\bar{dZ}}{dt}}}$$

5 where:

$\frac{\hat{dZ}}{dt}$ is the magnitude of the minimum time derivative

of Z

$\frac{\hat{dZ}_n}{dt}$ is the compressed time derivative of z

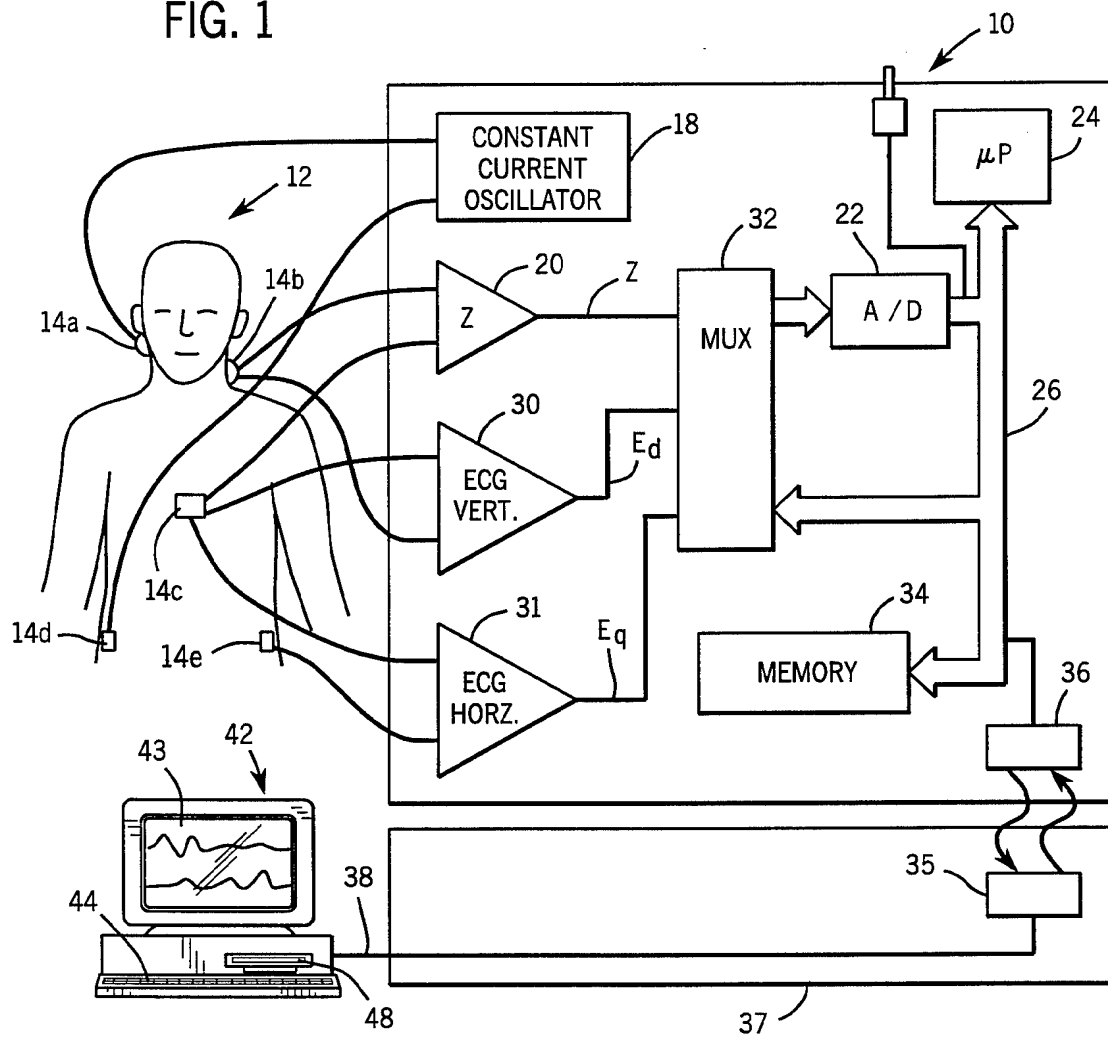
$\frac{\bar{dZ}}{dt}$ is the predetermined normal value and is an

10 average value of $\frac{\hat{dZ}}{dt}$ for a population.

16. The method of claim 14 wherein the predetermined normal value is substantially 1.73 ohms per second.

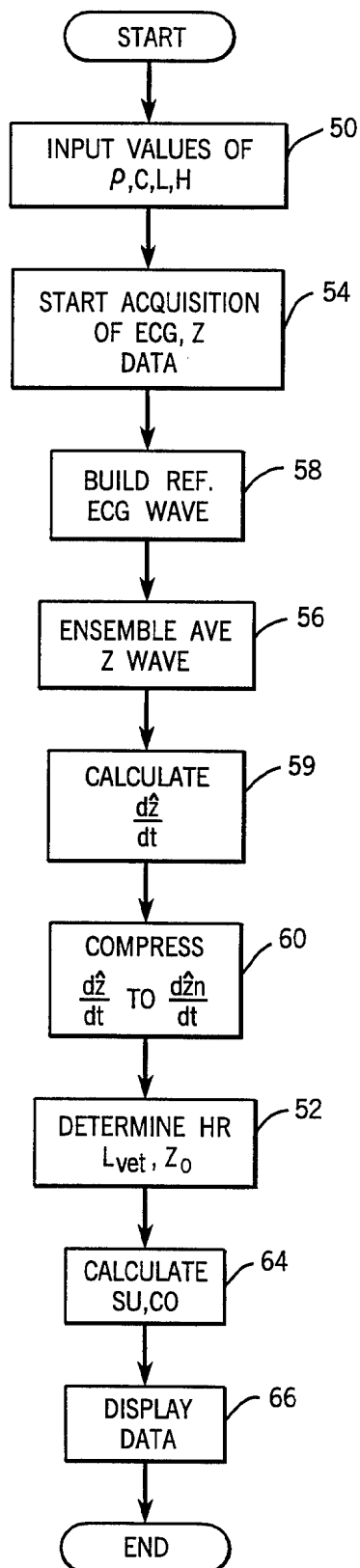
1 / 2

FIG. 1



2 / 2

FIG. 2



INTERNATIONAL SEARCH REPORT

Internat Application No

PCT/US 95/08856

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 A61B5/05 A61B5/029

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 6 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MEDICAL ELECTRONICS, vol. 2, April 1982 pages 93-97, BO SRAMEK 'Cardiac output by electrical impedance' cited in the application see the whole document ---	1,2,9,10
A	PROCEEDINGS OF THE NINTH ANNUAL CONFERENCE OF THE IEEE ENGINEERING IN MEDICINE AND BIOLOGY SOCIETY, vol. 3/4, November 1987 US, pages 1488-1489, QU ET AL. 'Portable impedance cardiograph for ambulatory subjects' see the whole document --- -/--	1-3,9-11

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

14 November 1995

Date of mailing of the international search report

23. 11. 95

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax (+31-70) 340-3016

Authorized officer

Chen, A

INTERNATIONAL SEARCH REPORT

Internat'l Application No
PCT/US 95/08856

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO,A,93 04627 (DREXEL UNIVERSITY) 18 March 1993 see page 50, line 5 - page 53, line 13 ---	1,2,9,10
A	MEDICAL & BIOLOGICAL ENGINEERING & COMPUTING, vol. 19, no. 5, September 1981 pages 638-644, MIYAMOTO ET AL. 'Continuous determination of cardiac output during exercise by the use of impedance plethysmography' see page 638, left column, line 25 - page 639, right column, line 31 -----	1-3,9-11